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William R. Schmidt, II
Wenderoth, Lind & Ponack, L.L.P.
1030 15th St., N.W.
Suite 400 East
Washington, DC 20005

In Re: Patent Term Extension
Application for
U.S. Patent No. 6,114,319

NOTICE OF FINAL DETERMINATION

A determination has been made that U.S. Patent No. 6,114,319, claims of which cover the human drug product DUREZOL® (difluprednate ophthalmic emulsion), is eligible for patent term extension under 35 U.S.C. § 156. The period of extension has been determined to be 371 days.

A single request for reconsideration of this final determination as to the length of extension of the term of the patent may be made if filed within one month of the date of this notice. Extensions of time under 37 CFR § 1.136(a) are not applicable to this time period. In the absence of a request for reconsideration, the Director will issue a certificate of extension, under seal, for a period of 371 days.

The period of extension has been calculated using the Food and Drug Administration determination of the length of the regulatory review period published in the Federal Register of July 29, 2009 (74 FR 37716). Under 35 U.S.C. § 156(c):

Period of Extension = RRP - PGRRP - DD - ½ (TP - PGTP)[1]
= 560- 0 - 0 - ½ (379 - 0)
= 371 days (1.0 years)

Since the regulatory review period began December 13, 2006, after the patent issued (September 5, 2000), the entire regulatory review period has been considered in the above determination of the length of the extension period 35 U.S.C. § 156(c). No determination of a lack of due diligence under 35 U.S.C. § 156(c)(1) was made.

Neither the limitations of 35 U.S.C. § 156(g)(6) nor 35 U.S.C. § 156(c)(3) operate to reduce the period of extension determined above.

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[1] Consistent with 35 U.S.C. § 156(c), "RRP" is the total number of days in the regulatory review period, "PGRRP" is the number of days of the RRP which were on and before the date on which the patent issued, "DD" is the number of days of the RRP that the applicant did not act with due diligence, "TP" is the testing phase period described in paragraphs (1)(B)(i), (2)(B)(i), (3)(B)(i), (4)(B)(i), and (5)(B)(i) of subsection (g) of 35 U.S.C. § 156, and "PGTP" is the number of days of the TP which were on and before the date on which the patent issued, wherein half days are ignored for purposes of the subtraction of ½ (TP - PGTP).

Upon issuance of the certificate of extension, the following information will be published in the Official Gazette:

U.S. Patent No.:	6,114,319
Granted:	September 5, 2000
Original Expiration Date[2]:	May 12, 2018
Applicant:	Masako Kimura et al.
Owner of Record:	Senju Pharmaceutical Co., Ltd.; Mitsubishi Chemical Corporation
Title:	Compositions Containing Difluprednate
Product Trade Name:	DUREZOL® (difluprednate ophthalmic emulsion)
Term Extended:	371 days
Expiration Date of Extension:	May 18, 2019

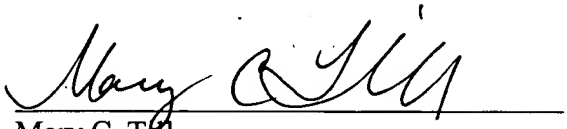
Any correspondence with respect to this matter should be addressed as follows:

By mail: Mail Stop Hatch-Waxman PTE
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450.

By FAX: (571) 273-7755

[2]Subject to the provisions of 35 U.S.C. § 41(b).

Telephone inquiries related to this determination should be directed to the undersigned at (571) 272-7755.



Mary C. Till
Legal Advisor
Office of Patent Legal Administration
Office of the Associate Commissioner
for Patent Examination Policy

cc: Office of Regulatory Policy
Food and Drug Administration
10903 New Hampshire Ave., Bldg. 51, Rm. 6222
Silver Spring, MD 20993-0002

Attention: Beverly Friedman

RE: DUREZOL® (difluprednate ophthalmic emulsion)
Docket No.: FDA-2009-E-0021